**Amendments to the Claims:** 

This listing of claims will replace all prior versions, and listings, of claims in the

application:

**Listing of Claims:** 

Claim 1 (withdrawn): A multiplexed capillary electrophoresis system for the

separation and detection of proteins and peptides, comprising:

(a) an array of coplanar parallel capillary electrophoresis tubes, each having a

first and a second end, said first ends being arranged in a two-dimensional

array having a spacing corresponding to that of an array of wells of a

microtiter plate;

(b) an apparatus arranged to selectively deliver sieving matrix and a selected one

of a plurality of liquids to said capillary tube second ends; and

(c) a scanning means for exciting and detecting radiation from said array of

capillary tubes.

Claim 2 (withdrawn): The system of claim 1 wherein said sieving matrix is a size

based sieving matrix.

Claim 3 (withdrawn): The system of claim 2 wherein said sieving matrix includes

dextran.

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Claim 4 (withdrawn): The system of claim 2 wherein said sieving matrix includes

galactomannans.

Claim 5 (withdrawn): A multiplexed capillary electrophoresis system for the

separation and detection of biomolecules, comprising:

(a) an array of coplanar parallel capillary electrophoresis tubes, each having a

first end and a second end, said first ends being arranged in a two-

dimensional array having a spacing corresponding to that of an array of wells

of a microtiter plate;

(b) an apparatus arranged to selectively deliver sieving matrix and a selected one

of a plurality of liquids to said capillary tube second end; and

(c) a scanning means for exciting and detecting endogenous fluorescence

radiation of the biomolecules from said array of capillary tubes.

Claim 6 (withdrawn): The system of claim 5 wherein said scanning means includes

a laser capable of producing radiation of an ultraviolet wavelength.

Claim 7 (withdrawn): The system of claim 6 wherein said laser is a multiplied

titanium sapphire laser.

Claim 8 (withdrawn): The system of claim 5 wherein said sieving matrix is a size

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based sieving matrix.

Claim 9 (withdrawn): The system of claim 8 wherein said sieving matrix includes dextran.

Claim 10 (withdrawn): The system of claim 8 wherein said sieving matrix includes galactomannans.

Claim 11 (withdrawn): The system of claims 1 wherein the array of coplanar parallel capillary electrophoresis tubes comprises at least 16 capillaries.

Claim 12 (withdrawn): The system of claims 1 wherein the array of coplanar parallel capillary electrophoresis tubes comprises at least 96 capillaries.

Claim 13 (withdrawn): The system of claims 1 wherein the array of coplanar parallel capillary electrophoresis tubes comprises at least 384 capillaries.

Claim 14 (currently amended): A method of separating and detecting <u>proteins</u> components-in a complex biological sample by two dimensional separations, comprising:

(a) subjecting said sample to a first separation and detection means to <u>separate</u>

<u>said sample into</u> a plurality of fractions;

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(b) collecting said plurality of fractions in a fraction collection means; and

(c) subjecting more than one fraction of said plurality of fractions

simultaneously to a second separation and detection means consisting of a

highly parallel capillary gel electrophoresis system which utilizes a size-

based sieving matrix, wherein said complex biological sample is a cell lysate

second separation and detection means is based on a different property of the

component being separated than said first separation and detection means.

Claim 15 (original): The method of claim 14, further comprising the step of dye

labeling said complex biological sample before subjecting said sample to the first

separation and detection means.

Claim 16 (original): The method of claim 14, further comprising the step of dye

labeling said fractions of the complex biological sample after collecting said

fractions into said fraction collection means.

Claim 17 (original): The method of claim 14, further comprising the step of

adding controls labeled with mobility-matched dyes to the fractions after said

collecting step.

Claim 18 (original): The method of claim 14, whereas the first separation and

detection means consists of HPLC, FPLC, ion exchange chromatography,

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hydrophobic interaction chromatography, affinity chromatography, isoelectric

focusing, isotachophoresis, capillary zone electrophoresis, micellar electrokinetic

chromatography, electrochromatography, field flow fractionation, solid phase

extraction, liquid phase extraction, or any other standard separation means.

Claim 19 (cancelled)

Claim 20 (currently amended): The method of claim 14 claim 19, wherein

galactomannans is used as the a-sieving matrix in the second separation and

detection means.

Claim 21 (currently amended): The method of claim 14-claim 19, wherein dextran

is used as the a-sieving matrix in the second separation and detection means.

Claim 22 (original): The method of claim 14, whereas said fraction collection

means consists of a microtiter plate.